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FILE COVERS 1907 - 11 Sep 2008 VOL 149 ISS 11 FILE LAST UPDATED: 10 Sep 2008 (20080910/ED)

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G1 X, Ak, CN, O

Structure attributes must be viewed using STN Express query preparation.

L2 727 SEA FILE=REGISTRY SSS FUL L1

L3 15 SEA FILE=CAPLUS L2

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L3 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:831355 CAPLUS

DOCUMENT NUMBER: 149:152831

TITLE: Process for preparation of 2-amino-5-cyanobenzamides

from the corresponding 5-halo compounds using metal cyanides in the presence of cuprous salts, iodides,

GΙ

and amines.

INVENTOR(S): Annis, Gary David; Bruening, Joerg; Currie, Martin

James; Dumas, Donald Joseph; Shapiro, Rafael E. I. Du Pont de Nemours and Company, USA

SOURCE: PCT Int. Appl., 63 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT ASSIGNEE(S):

PA	TENT	NO.			KIN	D	DATE			APPL	ICAT	ION I	NO.			ATE	
WC	2008	0825	02		A2	_	2008	0710		——— WO 2	 007-1	JS25	800			0071	
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		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FΙ,
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PRIORIT	Y APP	LN.	INFO	. :						US 2	006-	8763	94P	-	P 2	0061	221
										US 2	007-	9024	65P		P 2	0070	221
OTHER S	OURCE	(S):			CAS	REAC	T 14	9:15	2831	; MA	RPAT	149	:152	831			

$$R^2$$
 $NH_2$ 
 $NC$ 
 $COR^1$  I
 $Y$ 
 $COR^1$  II

Title compds. (I; R1 = NHR3, OR4; R2 = Me, C1; R3 = H, alkyl, cyclopropyl, cyclopropylcyclopropyl, cyclopropylmethyl, methylcyclopropyl; R4 = H, alkyl), were prepared by treatment of the corresponding halides (II; Y = Br, C1; R1, R2 as above) with a metal cyanide in the presence of a Cu(I) salt, an iodide salt, and R5HNCR6R7(CR8R9)nCR10R11XR12 (X = NR13, O; n = 0, 1; R5, R7, R8, R9, R11, R12 = H, alkyl; R6, R10 = H, alkyl, Ph; R13 = H, Me; n = 0, 1). Thus, 2-amino-5-bromo-N,3-dimethylbenzamide (preparation given), NaCN, CuI, and N,N'-dimethylethylenediamine were heated in xylenes at 140° for 4.5 h to give 2-amino-5-cyano-N,3-dimethylbenzamide.

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RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
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                                               1042429-14-0P
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                               1042429-50-4P
                                               1042429-52-6P
RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
preparation); PREP (Preparation)
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(prepn of aminocyanobenzamides from the corresponding halo compds. using metal cyanides in the presence of cuprous salts, iodides, and amines)

L3 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:735755 CAPLUS

DOCUMENT NUMBER: 149:47043

TITLE: Synergistic insecticidal compositions comprising an

anthranilamide derivative

INVENTOR(S): Koyanagi, Toru; Morita, Masayuki; Yoneda, Tetsuo;

Ueda, Tsuyoshi; Kiriyama, Kazuhisa; Hamamoto, Taku

PATENT ASSIGNEE(S): Ishihara Sangyo Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 82pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	TENT	NO.			KIN	D	DATE			APPL	ICAT	ION I	.OV		D	ATE	
WO	2008	 0727	83		A1	_	2008	0619	1	WO 2	007-	JP74:	 372		2	0071	212
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	ΒA,	BB,	BG,	BH,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FΙ,
		GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,
	KM, KN, K				KR,	KΖ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ΜE,
	MG, MK, MI					MX,	MY,	MZ,	NA,	NG,	NΙ,	NO,	NZ,	OM,	PG,	PH,	PL,
	MG, MK, MI PT, RO, RS				RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	ΤJ,	TM,	TN,
		TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW				
	RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FΙ,	FR,	GB,	GR,	HU,	ΙE,
		IS,	ΙT,	LT,	LU,	LV,	MC,	MT,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BW,
		GH,	GM,	ΚE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,
		BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM									
PRIORIT	PRIORITY APPLN. INFO.:									JP 2	006-	3365	85	Ž	A 2	0061	214
										JP 2	007-	1050	29	Ž	A 2	0070	412

OTHER SOURCE(S): MARPAT 149:47043

GΙ

$$R^{1?}$$
 $NH-CO$ 
 $N-N$ 
 $R^{2}$ 
 $R^{3}q$ 
 $O_{n}=N$ 

AB Synergistic insecticidal and ectoparasiticidal compns. contain a anthranilamide derivs. I [R1a, R1b = halo; R2, R3 = halo, alkyl, haloalkyl, alkoxy, haloalkoxy or cyano; A = alkyl substituted by Y; Y = (un)substituted C3-4 cycloalkyl; n = 0 or 1; q = 0, 1-4] and another pesticide.

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

TT 1031756-99-6 1031757-01-3 1031757-03-5 1031757-05-7 1031757-07-9 1031757-10-4 1031757-14-8 1031757-17-1 1031757-19-3 1031757-21-7 1031757-23-9 1031757-26-2 1031757-28-4 1031757-30-8 1031757-32-0

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1031757-34-2 1031757-36-4 1031757-38-6
    1031757-41-1 1031757-43-3 1031757-45-5
    1031757-47-7 1031757-49-9 1031757-51-3
    1031757-54-6 1031757-56-8 1031757-58-0
    1031757-60-4 1031757-62-6 1031757-64-8
    1031757-66-0 1031757-69-3 1031757-71-7
    1031757-74-0 1031757-76-2 1031757-78-4
    1031757-80-8 1031757-82-0 1031757-84-2
    1031757-86-4 1031757-88-6 1031757-90-0
    1031757-92-2 1031757-94-4
    RL: AGR (Agricultural use); BUU (Biological use, unclassified); BIOL
     (Biological study); USES (Uses)
        (synergistic insecticidal compn)
    112410-23-8D, Tebufenozide, mixts. with anthranilamide derivs.
ΤТ
    112839-32-4D, Furconazole-cis, mixts. with anthranilamide derivs.
    114369-43-6D, Fenbuconazole, mixts. with anthranilamide derivs.
    115852-48-7D, Fenoxanil, mixts. with anthranilamide derivs.
    116255-48-2D, Bromuconazole, mixts. with anthranilamide derivs.
    116714-46-6D, Novaluron, mixts. with anthranilamide derivs.
    117428-22-5D, Picoxystrobin, mixts. with anthranilamide derivs.
    118134-30-8D, Spiroxamine, mixts. with anthranilamide derivs.
    119168-77-3D, Tebufenpyrad, mixts. with anthranilamide derivs.
    119446-68-3D, DIfenoconazole, mixts. with anthranilamide derivs.
    119544-94-4D, Protrifenbute, mixts. with anthranilamide derivs.
    119791-41-2D, Emamectin, mixts. with anthranilamide derivs.
    120068-37-3D, Fipronil, mixts. with anthranilamide derivs.
                                                                  120116-88-3D,
    Cyazofamid, mixts. with anthranilamide derivs. 120928-09-8D, Fenazaquin,
    mixts. with anthranilamide derivs. 121451-02-3D, Noviflumuron, mixts.
    with anthranilamide derivs. 121552-61-2D, Cyprodinil, mixts. with
    anthranilamide derivs. 122453-73-0D, Chlorfenapyr, mixts. with
    anthranilamide derivs. 123312-89-0D, Pymetrozine, mixts. with
    anthranilamide derivs. 124495-18-7D, Quinoxyfen, mixts. with
    anthranilamide derivs. 125116-23-6D, Metconazole, mixts. with
    anthranilamide derivs. 125225-28-7D, Ipconazole, mixts. with
                            126069-54-3D, Phosphocarb, mixts. with
    anthranilamide derivs.
    anthranilamide derivs.
                             126833-17-8D, Fenhexamid, mixts. with
    anthranilamide derivs.
                             129558-76-5D, Tolfenpyrad, mixts. with
    anthranilamide derivs.
                             130000-40-7D, Thifluzamide, mixts. with
                             131341-86-1D, Fludioxonil, mixts. with
    anthranilamide derivs.
    anthranilamide derivs.
                             131807-57-3D, Famoxadone, mixts. with
    anthranilamide derivs.
                             131860-33-8D, Azoxystrobin, mixts. with
    anthranilamide derivs.
                             133855-98-8D, Epoxiconazole, mixts. with
                             134098-61-6D, Fenpyroximate, mixts. with
    anthranilamide derivs.
                             135410-20-7D, Acetamiprid, mixts. with
    anthranilamide derivs.
                             136426-54-5D, Fluquinconazole, mixts. with
    anthranilamide derivs.
                              138261-41-3D, Imidacloprid, mixts. with
    anthranilamide derivs.
                              139920-32-4D, Diclocymet, mixts. with
    anthranilamide derivs.
                              139968-49-3D, Metaflumizone, mixts. with
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                              140163-89-9D, Imicyafos, mixts. with
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                              140923-17-7D, Iprovalicarb, mixts. with 141517-21-7D, Trifloxystrobin, mixts. with
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                              143390-89-0D, Kresoximmethyl, mixts. with
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                              143807-66-3D, Chromafenozide, mixts. with
                              148477-71-8D, Spirodiclofen, mixts. with
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                              149508-90-7D, Sipconazole, mixts. with
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                              149877-41-8D, Bifenazate, mixts. with
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                              149961-52-4D, Dimoxystrobin, mixts. with
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    anthranilamide derivs. 153233-91-1D, Etoxazole, mixts. with
                             153719-23-4D, Thiamethoxam, mixts. with
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155569-91-8D, Emamectin benzoate, mixts. with
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anthranilamide derivs. 156052-68-5D, Zoxamide, mixts. with
anthranilamide derivs. 158062-67-0D, Flonicamid, mixts. with
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anthranilamide derivs. 161326-34-7D, Fenamidone, mixts. with
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anthranilamide derivs. 168316-95-8D, Spinosad, mixts. with
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                188425-85-6D, Boscalid, mixts. with anthranilamide derivs.
189278-12-4D, Proquinazid, mixts. with anthranilamide derivs.
203313-25-1D, Spirotetramat, mixts. with anthranilamide derivs.
210880-92-5D, Clothianidin, mixts. with anthranilamide derivs.
211867-47-9D, Flumorph, mixts. with anthranilamide derivs. 220899-03-6D,
Metrafenone, mixts. with anthranilamide derivs. 223419-20-3D,
Profluthrin, mixts. with anthranilamide derivs.
                                                                                223580-51-6D, Tiadinil,
mixts. with anthranilamide derivs. 229977-93-9D, Fluacrypyrim, mixts.
with anthranilamide derivs. 240494-70-6D, ,Metofluthrin, mixts. with
anthranilamide derivs. 248593-16-0D, Orysastrobin, mixts. with
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anthranilamide derivs. 315208-17-4D, Pyrafluprole, mixts. with
anthranilamide derivs. 337458-27-2D, Pyrifluquinazon, mixts. with
anthranilamide derivs. 348635-87-0D, Amisulbrom, mixts. with
anthranilamide derivs. 361377-29-9D, Fluoxastrobin, mixts. with
anthranilamide derivs. 394730-71-3D, Pyriprole, mixts. with
anthranilamide derivs. 400882-07-7D, Cyflumetofen, mixts. with
anthranilamide derivs. 413615-35-7D, Benthiavalicarb-, mixts. with
anthranilamide derivs. 500008-45-7D, Chlorantraniliprole, mixts. with
anthranilamide derivs. 560121-52-0D, Cyenopyrafen, mixts. with
anthranilamide derivs. 863549-51-3D, Lepimectin, mixts. with
anthranilamide derivs. 935545-74-7D, DE 175, mixts. with anthranilamide
derivs. 937279-54-4D, HGW 86, mixts. with anthranilamide derivs.
946494-18-4D, UBF 307, mixts. with anthranilamide derivs. 946494-19-5D,
KIF 7767, mixts. with anthranilamide derivs.
                                                                              946494-20-8D, Syngenta
446510, mixts. with anthranilamide derivs. 1031756-98-5D, mixts.
containing 1031757-98-8D, mixts. containing 1031758-00-5D,
                               1032110-39-6D, BCM 062, mixts. with anthranilamide derivs.
mixts. containing
1032111-18-4D, BCM 061, mixts. with anthranilamide derivs.
1032111-42-4D, BCF 051, mixts. with anthranilamide derivs.
RL: AGR (Agricultural use); BUU (Biological use, unclassified); BIOL
 (Biological study); USES (Uses)
      (synergistic insecticidal compns.)
ANSWER 3 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
```

2008:734496 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 149:79593

TITLE: Process for production of anthranilamide compound INVENTOR(S): Koyanagi, Toru; Yamamoto, Kazuhiro; Yoneda, Tetsuo; Kanbayashi, Shigehisa; Tanimura, Toyoshi; Taguchi,

GΙ

Yohei; Yoshida, Tatsunori

PATENT ASSIGNEE(S): Ishihara Sangyo Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 95pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

	PATENI	NO.			KIN	D	DATE		-	APPL	ICAT	ION I	NO.		D	ATE	
	WO 200	80727	45		A1	_	2008	0619		WO 2	007-	JP74	 169		2	0071	214
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BH,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FI,
		GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,
		KM,	KN,	KP,	KR,	KΖ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ΜE,
		MG,	MK,	MN,	MW,	MX,	MY,	MZ,	NA,	NG,	NΙ,	NO,	NZ,	OM,	PG,	PH,	PL,
		PT,	RO,	RS,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	ΤJ,	TM,	TN,
		TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW				
	RV	: AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,
		IS,	ΙΤ,	LT,	LU,	LV,	MC,	MT,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	ΤG,	BW,
		GH,	GM,	KΕ,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	ΑM,	ΑZ,
		BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM									
PRIOR	ITY AE	PLN.	INFO	.:					1	JP 2	006-	3391	00		A 2	0061	215
									1	JP 2	007-	1527	18	1	A 2	0070	608
OTHER	SOURC		MAR:	PAT	149:	7959.	3										

Disclosed is a process for producing a specific anthranilamide compound or a salt thereof. Specifically disclosed is a process for producing an anthranilamide compound represented by the formula I: [wherein R1a and R3 independently represent a halogen or a haloalky1; R2 represents a cyclopropylalky1 or a cyclobutylalky1; and Hal represents a chlorine atom or a bromine atom] or a salt thereof, which comprises the step of selectively halogenating a compound represented by the formula I: [wherein R1a, R2 and R3 are as defined above, Hal = H]. For example, II was provided in a multi-step synthesis starting from the reaction of Et 2-furoylpyruvate with 3-chloro-2-hydrazylpyridine.

REFERENCE COUNT: 55 THERE ARE 55 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

IT 1621-24-5P, (1-Cyclopropylethyl)amine 51761-72-9P, Cyclopropylmethylketoxime 107855-32-3P 112881-69-3P 112881-76-2P

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1033344-64-7P, 1-(3-Chloropyridin-2-y1)-5-(2-fury1)-1H-pyrazole-3-
    carboxylic acid 1033344-84-1P, 3-[(Benzyloxycarbonyl)amino]-1-(3-
    chloropyridin-2-yl)-1H-5-pyrazolecarboxylic acid 1033344-87-4P,
    3-[(Benzyloxycarbonyl)amino]-1-(3-chloropyridin-2-yl)-5-(2-furyl)-1H-
    pyrazole 1033407-53-2P
                              1033407-55-4P 1033407-56-5P,
    N-[4-Chloro-2-(1-cyclopropylethylcarbamoyl)phenyl]-1-(3-chloropyridin-2-
    v1)-3-hydroxy-4,5-dihydro-1H-pyrazole-5-carboxamide 1033407-57-6P
    1033407-58-7P, 3-Bromo-N-[4-chloro-2-(1-cyclopropylethylcarbamoyl)phenyl]-
    1-(3-chloropyridin-2-yl)-4,5-dihydro-1H-pyrazole-5-carboxamide
    1033407-59-8P, 3-Bromo-N-[4-chloro-2-(1-
    cyclopropylethylcarbamoyl)phenyl]-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxamide 1033407-60-1P
                                 1033407-61-2P, 2-Amino-5-chloro-N-(1-
    cyclopropylethyl)benzamide 1033407-63-4P 1033407-64-5P, Pentyl
    3-bromo-1-(3-chloropyridin-2-yl)-4,5-dihydro-1H-pyrazole-5-carboxylate
    1033407-65-6P, Pentyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxylate 1033407-66-7P, Phenyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-
    pyrazole-5-carboxylate 1033407-67-8P, 2-Amino-3-bromo-5-chloro-N-(1-
    cyclopropylethyl)benzamide 1033407-69-0P, Ethyl 1-(3-chloropyridin-2-yl)-
    5-(2-furyl)-4,5-dihydro-1H-pyrazole-3-carboxylate 1033407-70-3P,
    3-[(Benzyloxycarbonyl)amino]-1-(3-chloropyridin-2-yl)-1H-5-
    pyrazolecarboxylic acid phenyl ester 1033407-72-5P, 3-Amino-1-(3-
    chloropyridin-2-yl)-1H-5-pyrazolecarboxylic acid phenyl ester
    1033407-73-6P, Benzyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxylate 1033407-75-8P 1033407-76-9P
                                                 1033407-78-1P,
     5-Chloro-N-(1-cyclopropylethyl)-2-nitrobenzamide
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation of anthranilamide compound)
ΙT
    1031756-98-5P, 3-Bromo-N-[2-bromo-4-chloro-6-[[(1-
    cyclopropylethyl)amino]carbonyl]phenyl]-1-(3-chloropyridin-2-yl)-1H-
    pyrazole-5-carboxamide
                            1033407-74-7P, 4-Methoxybenzyl
    3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-carboxylate
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of anthranilamide compound)
    ANSWER 4 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        2008:733566 CAPLUS
DOCUMENT NUMBER:
                        149:79590
TITLE:
                        Process for preparation of anthranilamide compound by
                        using novel pyrazole compound as intermediate
INVENTOR(S):
                        Koyanagi, Toru; Hisamatsu, Akihiro
PATENT ASSIGNEE(S):
                        Ishihara Sangyo Kaisha, Ltd., Japan
                        PCT Int. Appl., 59pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        Japanese
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:
    PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO. DATE
                                         WO 2007-JP74166 20071214
                              20080619
    WO 2008072743
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        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA,
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            GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG,
            KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME,
            MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL,
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PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN,

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TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.: JP 2006-339100 A 20061215

JP 2007-128991 A 20070515 JP 2007-137551 A 20070524

OTHER SOURCE(S): MARPAT 149:79590

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AB Disclosed is a method for producing an anthranilamide compound or a salt thereof. Specifically disclosed is a method for producing an anthranilamide compound represented by the formula I [R1 = halo, alkyl, alkenyl, etc.; A = (un)substituted alkyl; X = halo; m = 0-4] or a salt thereof, which is characterized in that a compound represented by the formula I [R1, A and m are defined as above; X = NH2] is diazotized and then reacted with copper halide, copper metal or an alkyl halide. For example, II was provided in a multi-step synthesis starting from the reaction of Et 2-furoylpyruvate with 3-chloro-2-hydrazylpyridine.

REFERENCE COUNT: 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS

REFERENCE COUNT: 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

IT 1031756-98-5P

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(preparation of anthranilamide compound by using novel pyrazole compound as intermediate)

L3 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:703045 CAPLUS

DOCUMENT NUMBER: 149:53719

TITLE: Process for preparing 2-amino-5-cyanobenzoates by

treatment of 2-amino-5-halobenzoates with alkali metal

nitriles in the presence of palladium phosphine

catalysts.

INVENTOR(S): Bruening, Joerg; Casalnuovo, Albert Loren; Grushin,

Vladimir

PATENT ASSIGNEE(S): E. I. du Pont de Nemours and Company, USA

SOURCE: PCT Int. Appl., 50pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

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## PATENT INFORMATION:

	PAI	ENT :	NO.			KIN	D	DATE			APPL	ICAT	ION 1	NO.		D.	ATE	
	WO	2008	0701	 58		A1	_	2008	0612		WO 2	007-	US25	005		2	0071	205
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PRIO:	RIORITY APPLN. INFO.:										US 2	006-	8730	58P		P 2	0061	206
OTHE	THER SOURCE(S):						REAC	T 14	9:53	719;	MAR	PAT	149:	5371	9			
GT																		

$$R^2$$
 $NH_2$ 
 $NC$ 
 $COR^1$  I
 $R^2$ 
 $NH_2$ 
 $NH_2$ 
 $COR^1$  II

AΒ Title compds. (I; R1 = NHR3, OR4; R2 = Me, C1; R3 = H, alkyl, cyclopropyl, cyclopropylmethyl, methylcyclopropyl; R4 = H, alkyl; with a proviso); were prepared by treatment of (II; X = Br, Cl; other variables as above) with M1CN (M1 = alkali metal) in the presence of ≥1 ether and nitrile solvent, ≥1 palladium tertiary phosphine catalyst. Thus, 2-amino-5-bromo-N,3-dimethylbenzamide (preparation given), Zn, NaCN, and a catalyst solution prepared from tris(dibenzylideneacetone)dipalladium and tri-tert-butylphosphine were stirred together in THF for 63 h at 25° to give >99% conversion to 2-amino-5-cyano-N,3-

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dimethylbenzamide.
REFERENCE COUNT:
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    RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
    preparation); PREP (Preparation)
        (preparation of aminocyanobenzoates by treatment of aminohalobenzoates with
        alkali metal nitriles in the presence of palladium phosphine catalysts)
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(preparation of aminocyanobenzoates by treatment of aminohalobenzoates with alkali metal nitriles in the presence of palladium phosphine catalysts)

L3 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:673452 CAPLUS

DOCUMENT NUMBER: 149:10005

TITLE: Preparation of (heterocyclyl) N-

cyanoalkylanthranilamides as insecticides and

acaricides

INVENTOR(S): Muehlebach, Michel; Craig, Gerald Wayne

PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.

SOURCE: PCT Int. Appl., 91pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PAI	PATENT NO.					D	DATE		1	APPL	ICAT	ION I	. O <i>V</i>		D.	ATE	
	WO	2008	0648	 91		A1	_	2008	0605	1	WO 2	 007-1	==== EP10:	 370		2	0071	129
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OTHER SOURCE(S): MARPAT 149:10005

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AB Title compds. [I; D = (substituted) Ph, pyridyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidyl; n = 0-3; R1 = halo, OH, NO2, alkyl, alkenyl, alkynyl,cycloalkyl, alkylthio, cycloalkylamino, (substituted) Ph, PhCH2, PhO, etc.; R2, R3 = H, (substituted) alkyl, alkenyl, alkynyl, cycloalkyl; E1, E2 = 0, S; R4 = (substituted) alkyl, cycloalkyl; R5 = (substituted) cycloalkyl, cycloalkylalkyl], were prepared Thus, title compound (II) was prepared in 4 steps from 2-amino-5-chloro-3-methylbenzoic acid, cyclopropyl Me ketone, and 2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazole-3carbonyl chloride. II and other I at 200 ppm gave >80% control of Spodoptera littoralis. REFERENCE COUNT: THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT 1034824-68-4P 1034824-71-9P 1034824-74-2P 1034824-77-5P ΤT 1034824-80-0P 1034824-82-2P 1034824-84-4P 1034824-87-7P 1034824-93-5P 1034824-96-8P 1034824-99-1P 1034824-90-2P 1034825-02-9P 1034825-05-2P 1034825-08-5P 1034825-11-0P 1034825-14-3P 1034825-17-6P 1034825-20-1P 1034825-23-4P 1034825-26-7P 1034825-29-0P 1034825-32-5P 1034825-35-8P 1034825-41-6P 1034825-38-1P 1034825-44-9P 1034825-47-2P 1034825-56-3P 1034825-50-7P 1034825-53-0P 1034825-59-6P 1034825-68-7P 1034825-62-1P 1034825-65-4P 1034825-72-3P 1034825-81-4P 1034825-75-6P 1034825-78-9P 1034825-84-7P 1034825-93-8P 1034825-87-0P 1034825-90-5P 1034825-96-1P 1034825-99-4P 1034826-02-2P 1034826-05-5P 1034826-08-8P 1034826-11-3P 1034826-14-6P 1034826-17-9P 1034826-20-4P 1034826-23-7P 1034826-26-0P 1034826-29-3P 1034826-32-8P 1034826-35-1P 1034826-38-4P 1034826-41-9P 1034826-44-2P 1034826-47-5P 1034826-50-0P 1034826-52-2P 1034826-54-4P 1034826-56-6P 1034826-59-9P 1034826-62-4P 1034826-64-6P 1034826-67-9P 1034826-70-4P 1034826-73-7P 1034826-75-9P 1034826-78-2P 1034826-81-7P 1034826-84-0P 1034826-86-2P 1034826-89-5P 1034826-92-0P 1034826-95-3P 1034826-97-5P 1034826-99-7P 1034827-01-4P 1034827-04-7P 1034827-07-0P 1034827-09-2P 1034827-11-6P 1034827-13-8P 1034827-15-0P 1034827-17-2P 1034827-19-4P 1034827-23-0P 1034827-25-2P 1034827-27-4P 1034827-29-6P 1034827-31-0P 1034827-33-2P 1034827-36-5P 1034827-38-7P 1034827-41-2P 1034827-44-5P 1034827-47-8P 1034827-49-0P 1034827-51-4P 1034827-53-6P 1034827-55-8P 1034827-58-1P 1034827-60-5P 1034827-62-7P 1034827-64-9P 1034827-66-1P 1034827-69-4P 1034827-71-8P 1034827-73-0P 1034827-77-4P 1034827-79-6P 1034827-75-2P 1034827-81-0P 1034827-85-4P 1034827-87-6P 1034827-83-2P 1034827-89-8P 1034827-91-2P 1034827-93-4P 1034827-95-6P 1034827-97-8P 1034827-99-0P 1034828-01-7P 1034828-03-9P 1034828-05-1P 1034828-07-3P 1034828-09-5P 1034828-11-9P 1034828-13-1P 1034828-15-3P 1034828-17-5P 1034828-19-7P 1034828-21-1P 1034828-23-3P 1034828-25-5P 1034828-27-7P 1034828-29-9P 1034828-31-3P 1034828-33-5P 1034828-35-7P 1034828-37-9P 1034828-39-1P 1034828-41-5P 1034828-43-7P 1034828-45-9P 1034828-47-1P 1034828-53-9P 1034828-49-3P 1034828-51-7P 1034828-55-1P 1034828-57-3P 1034828-59-5P 1034828-61-9P 1034828-64-2P

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     (Prophetic); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (preparation of (heterocyclyl) N-cyanoalkylanthranilamides as insecticides
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RL: AGR (Agricultural use); BSU (Biological study, unclassified); PRPH
(Prophetic); SPN (Synthetic preparation); BIOL (Biological study); PREP
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RL: AGR (Agricultural use); BSU (Biological study, unclassified); PRPH
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    ANSWER 7 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         2008:221501 CAPLUS
                         148:231873
DOCUMENT NUMBER:
TITLE:
                         Crop vigor and yield enhancement and
                         arthropod-vectored plant disease disruption by
                         carboxamide derivatives
INVENTOR(S):
                         Annan, Isaac Billy; Marcon, Paula Cristina Rodrigues
                         Gouveia; Portillo, Hector Eduardo
PATENT ASSIGNEE(S):
                         E. I. Du Pont de Nemours and Company, USA
SOURCE:
                         PCT Int. Appl., 23pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
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GΙ

PA	TENT		KIN	D	DATE			APPL	ICAT	ION I	NO.		D.	ATE			
WO	2008	0211	 52		A2	_	2008	0221		WO 2	 007-1	US17	 673		2	0070	808
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PRIORIT	PRIORITY APPLN. INFO.:									US 2	006-	8368	92P		P 2	0060	809
										US 2	006-	8370	59P		P 2	0060	810
										US 2	006-	8504	54P		P 2	0061	010
OTHER S	OURCE		MAR:	PAT	148:	2318	73										

AΒ The anthranilamide anthropodicides I (X = N, CF, CCl, CBr or CI; R1 = Me, Cl, Br or F; R2 = H, F, Cl, Br or CN; R3 = F, Cl, Br, C1-4 haloalkyl or haloalkoxy; R4a = H, C1-4 alkyl cyclopropylmethyl or 1-cyclopropylethyl; R4b = H or Me; R5, r6 = H, F, C1 or Br) or their N-oxides, as well as phthalic acid derivs. (Markush given) enhance crop vigor and crop yield and disrupt infectious disease transmission by arthropod pests. 272451-65-7 ΙT 438450-41-0 500008-00-4 500008-44-6 500008-45-7 736994-60-8 736994-63-1 736995-23-6 871238-02-7 871238-04-9 882401-50-5 886583-54-6 871238-03-8 886583-69-3 RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses) (crop vigor and yield enhancement and arthropod-vectored plant disease disruption by)

L3 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

Ι

ACCESSION NUMBER: 2008:91157 CAPLUS

DOCUMENT NUMBER: 148:191926

TITLE: Process for making 3-substituted 2-amino-5-

halobenzamides

INVENTOR(S): Davis, Richard Frank; Shapiro, Rafael; Taylor, Eric

Deguyon

PATENT ASSIGNEE(S): E. I. du Pont de Nemours and Company, USA

SOURCE: PCT Int. Appl., 38pp.

CODEN: PIXXD2

DOCUMENT TYPE: Pat.ent. English LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	ATENT 1	NO.			KIN	D	DATE			APPL	ICAT	ION 1	NO.		D.	ATE	
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PRIORIT	RIORITY APPLN. INFO.:					·	,			US 2	006-	8317	81P		P 2	0060	719
OTHER S	HER SOURCE(S):					REAC	T 14	8:19	1926	; MA	RPAT	148	:191	926			

## \* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

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AB
    Disclosed is a method for preparing I (R1 = H, alkyl, cyclopropyl,
    cyclopropylmethyl, or methylcyclopropyl; R2 = Me or C1; X = C1 or Br) by
    ring opening of II with R1-NH2 in the presence of a carboxylic acid and a
    method for preparing II by cyclization of III (R3 = (un)substituted alkyl or
    alkenyl) with phosphorus tribromide. Also disclosed is a method for
    preparing known insecticides IV (R4 = C1, Br, CF3, OCF2H or OCH2CF3; Z = CR7
    or N; R5 = F, C1 or Br; R6 = H, F or C1; R7 = H, F, C1 or Br) from I.
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RL: PRPH (Prophetic); SPN (Synthetic preparation); PREP (Preparation) (preparation of aminoarylcarboxamides via PBr3 induced cyclization of carboxamidobenzoic acids followed by ring opening with alkyl amines)

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L3 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
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ACCESSION NUMBER: 2007:1278752 CAPLUS

DOCUMENT NUMBER: 147:481495

TITLE: Use of carboxamide derivatives for disrupting the

reproductive performance of arthropods

INVENTOR(S): Annan, Isaac Billy; Flexner, John Lindsey; Marcon,

Paula Cristina Rodrigues Gouveia; Portillo, Hector

Eduardo

PATENT ASSIGNEE(S): E. I. Du Pont De Nemours and Company, USA

SOURCE: PCT Int. Appl., 34pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	TENT	NO.			KIN	D	DATE			APPL	ICAT	ION I	NO.		D	ATE	
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WO	2007	1266	36		A2		2007	1108		WO 2	007-	US69.	29		2	0070	320
WO	2007	1266	36		А3		2008	0320									
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GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA AU 2007243792 A1 20071108 AU 2007-243792 20070320 US 2006-787753P P 20060331 WO 2007-US6929 W 20070320 PRIORITY APPLN. INFO.: MARPAT 147:481495 OTHER SOURCE(S): GΙ

The carboxamide derivs. I (X = n, CF, CCl, CBr or CI; R1 = Me, Cl, Br or AΒ F; R2 = H, F, C1, Br or CN; R3 = F, C1, Br, haloalkyl or haloalkoxy; R4a = H, alkyl, cyclopropylmethyl or 1-cyclopropylethyl; R4b = H or Me; R5, R6 = H,F, Cl or Br), their N-oxides and salts.  $272451 - 65 - 7 \qquad 438450 - 41 - 0 \qquad 500008 - 00 - 4 \qquad 500008 - 44 - 6 \qquad 500008 - 45 - 7$ 736994-60-8 736994-63-1 736995-23-6 871238-02-7 871238-03-8 871238-04-9 882401-50-5 886583-54-6 886583-69-3 RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses) (use of carboxamide derivs. for disrupting the reproductive performance

ANSWER 10 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

Ι

ACCESSION NUMBER: 2006:1120573 CAPLUS

DOCUMENT NUMBER: 145:455006

of arthropods)

TITLE: Preparation of cyanoanthranilamides as insecticides

and acaricides

INVENTOR(S): Jeanguenat, Andre; O'Sullivan, Anthony; Muehlebach,

Michel; Trah, Stephan; Hall, Roger Graham

PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.

SOURCE: PCT Int. Appl., 100pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

KIND APPLICATION NO. PATENT NO. DATE DATE \_\_\_\_\_

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PRIORITY APPLN. INFO.:
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OTHER SOURCE(S): MARPAT 145:455006 GI

 $A_q X_p Y B$ 

Title compds. [I; E, Z = O, S; A = (substituted) alkylene, alkenylene, alkynylene, bivalent mono- or bicyclic ring; X = O, NH, alkylimino; Y = (substituted) mono- or bicyclic ring; p, q = 0, 1; B = (substituted) 3-4 membered (heterocyclic) ring; R1 = halo, NO2, cyano, OH, alkyl, alkenyl, alkynyl, cycloalkyl, haloalkyl, (substituted) Ph, PhCH2, PhO, etc.; n = 0-3; R2, R3 = H, alkyl, alkenyl, alkynyl, substituted cycloalkyl; D = (substituted) Ph, pyridyl, pyrrolyl, pyrazolyl, pyrimidyl], were prepared Thus, 2-[2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazol-3-yl]-8-methyl-4-oxo-4H-benzo[d][1,3]oxazine-6-carbonitrile, bicycloprop-1-ylamine hydrochloride (preparation given), and Et3N were heated together in THF at 60° for 8 h to give 2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [2-(bicycloprop-1-ylcarbamoyl)-4-cyano-6-methylphenyl]amide. The latter at 400 ppm showed >80% activity against Cydia pomonella.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT IT 1042422-55-8 1042425-11-5 1042425-51-3 1042426-12-9 1042426-58-3

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RL: PRPH (Prophetic)
   (Preparation of cyanoanthranilamides as insecticides and acaricides)
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L3 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:558556 CAPLUS

DOCUMENT NUMBER: 145:62886

TITLE: Anthranilamide derivatives as insecticides, and their preparation, pesticidal compositions and formulation Jeanguenat, Andre; O'Sullivan, Anthony Cornelius Syngenta Participations A.-G., Switz.

SOURCE: PCT Int. Appl., 136 pp.
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CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

P	ATENT	NO.			KIN		DATE				LICAT				D	ATE	
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											2005-				W 2	0051	207
OTHER S	SOURC	E(S):			CAS	REAC	T 14	5 <b>:</b> 62	886;	MA	RPAT	145:	6288	6			

## \* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

Compds. of formula I, and the agrochem. acceptable salts and all stereoisomers and tautomeric forms of the compds. of formula I can be used as agrochem. active ingredients and can be prepared in a manner known per se. Several examples on formulation of compds. of formula I is also disclosed in this invention. Compds. of formula I wherein E1 and W2 are independently O or S; R1 is halo, CN, NO2, OH, C1-6 (halo)alkyl, C2-6 (halo)alkenyl, C2-6 (halo)alkynyl, C3-6 (halo)cycloalkyl, C1-4 (halo)alkoxy, C1-4 (halo)alkylthio, C1-4 (halo)alkylsulfinyl, C1-4 (halo)alkylsulfonyl, C1-4 alkylamino, C2-4 dialkylamino, C3-6 cycloalkylamino, etc.; n is 0, 1, 2, 3, or 4; R2 and R3 are independently H, (un) substituted C1-6 alkyl, (un) substituted C2-6 alkenyl, (un) substituted C2-6 alkynyl, or (un) substituted C3-6 cycloalkyl; D is (un) substituted Ph, (un) substituted pyridyl, (un) substituted pyrazole, (un) substituted pyrrole, or (un) substituted pyrimidine; Y1a and Y2 are independently (un) substituted C1-6 alkylene, (un) substituted C2-6alkenylene, or (un)substituted C3-6 alkynylene, etc.; G is a bond, O, N-Z1, S or G1-C(=G2)-G3; G1 and G3 are independently a bond, O, S, or NZ2; G2 is O, S or NZ3; Z and Z1-Z3 are independently H, C1-6 (halo)alkyl, C2-6 (halo)alkenyl, C2-6 (halo)alkynyl, C3-6 (halo)cycloalkyl, C1-4 (halo)alkoxy, C1-4 (halo)alkylthio, etc.; Y3 is H, halo or C1-6

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(halo)alkyl; Y1b is a bond, or (un)substituted C1-6 alkylene,
     (un) substituted C2-6 alkenylene, or (un) substituted C3-6 alkynylene; and
     their tautomers, agrochem. utilizable salts and auxiliary are claimed.
     Example compound II was prepared by amidation of 6-\text{chloro}-2-[2-(3-\text{chloro})]
     chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazol-3-yl]-8-
     methylbenzo[d][1,3]oxazin-4-one with 1-amino-2-propanol; the resulting
     2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazol-3-carboxylic acid
     [4-chloro-2-(2-hydroxypropylcarbamoyl)-6-methylphenyl]amide underwent
     substitution with thioacetic acid to give thioacetic acid
     S-[2-(5-chloro-2-[2-(3-chloropyridin-2-y1)-5-trifluoromethyl-2H-pyrazole-
     3-carbonyl]amino]-3-methylbenzoylamino)-1-methylethyl] ester, which
     underwent deacetylation and methylation to give the corresponding Me thio
     ether, which underwent oxidation to give the corresponding sulfoxide, which
     reacted with trifluoroacetamide to give the corresponding
     N-trifluoroacetylated sulfoximide, which underwent deacetylation to give
     compound II. All the invention compds. were evaluated for their
     insecticidal activity. Some of the tested compds. showed good activity
     against Aphis craccivora, Diabrotica balteata, Heliothis virescens
     (application to foliar and egg), Myzus persicae (foliar and systemic
     application), Plutella xylostella and Spodoptera littoralis.
                               THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
                                                  891195-07-6P 891195-08-7P
891195-12-3P 891195-13-4P
     891195-04-3P
                  891195-05-4P
                                   891195-06-5P
ΙT
     891195-09-8P
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                                   891195-11-2P
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     891195-38-3P
     RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
        (agrochem. candidate; preparation of anthranilamide derivs. as insecticides)
     ANSWER 12 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         2006:496102 CAPLUS
DOCUMENT NUMBER:
                         144:462625
TITLE:
                         Preparation of anthranilamide derivative insecticides
                         and acaricides
INVENTOR(S):
                         Lahm, George Philip; Selby, Thomas Paul; Stevenson,
                         Thomas Martin; Taggi, Andrew Edmund; Bereznak, James
                         Francis
                        E.I. Dupont De Nemours and Co., USA
PATENT ASSIGNEE(S):
                         PCT Int. Appl., 97 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
                    KIND DATE
     PATENT NO.
                                        APPLICATION NO. DATE
                         A2 20060526 WO 2005-US42196
A3 20061221
                                             _____
                     A2
     WO 2006055922
                                                                    20051118
     WO 2006055922
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             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX,
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MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE,

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SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,
             VN, YU, ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
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                                                                    20050610
                                             WO 2005-US42196
                                                                 W 20051118
OTHER SOURCE(S):
                         MARPAT 144:462625
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AΒ

ΤТ

886583-68-2

N-oxides, and salts [J = (un)substituted Ph or <math>N-containing heterocyclyl; R1 =alkyl alkenyl, alkynyl, etc.; R2 = alkylcarbonyl, alkoxycarbonyl or (di)alkylaminocarbonyl; R3 = (cyclo)alkyl, alkenyl, alkynyl, alkoxy, etc. ; R4 = (un)substituted alkylcycloalkyl, alkenylcycloalkyl, alkynylcycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkylalkynyl, cycloalkenylalkyl or alkylcycloalkenyl, oxiranylalkyl, thiiranylalkyl, oxetanylalkyl, thietanylalkyl, 3-oxetanyl or 3-thietanyl; R5 = (cyclo)alkyl, haloalkyl, alkenyl alkynyl, etc.] are prepared as pesticides for controlling invertebrate pests, specifically insecticides and acaricides. 736995-23-6P 882401-50-5P 886583-28-4P ΙT 886583-29-5P 886583-30-8P 886583-31-9P 886583-32-0P 886583-33-1P 886583-34-2P 886583-35-3P 886583-36-4P 886583-37-5P 886583-38-6P 886583-39-7P 886583-40-0P 886583-41-1P 886583-42-2P 886583-43-3P 886583-44-4P 886583-45-5P 886583-46-6P 886583-47-7P 886583-48-8P 886583-49-9P 886583-50-2P 886583-51-3P 886583-52-4P 886583-53-5P 886583-54-6P 886583-55-7P 886583-56-8P 886583-57-9P 886583-58-0P 886583-59-1P 886583-69-3P RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation as insecticide and acaricides)

The anthranilamide derivs. I and their geometric and stereoisomers,

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)

886583-65-9 886583-66-0 886583-67-1

## (synergistic insecticide and acaricide)

L3 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:367128 CAPLUS

DOCUMENT NUMBER: 144:364548

Preparation of anthranilamide derivative acaricides TITLE:

and insecticides

INVENTOR(S): O'Sullivan, Anthony Cornelius; Hughes, Dave;

Jeanguenat, Andre; Muehlebach, Michel; Loiseleur,

PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.; Syngenta Limited

SOURCE: PCT Int. Appl., 152 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	TENT	NO.			KIN	D	DATE			APP	LICAT	ION	NO.		D	ATE	
	2006						2006 2006			WO 2	2005-	 EP10	 891		2	0051	010
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OTHER S	OURCE	(S):			CAS:	KEAC	T 14	4:36	4548	; M	ARPAT	144	:364	548			

SOURCE(S): CASREACT 144:364548; M

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The anthranilamides I [E, Z = O \text{ or } S; A, Y = alkylene, alkenylene,
AΒ
    alkynylene, etc.; X = O, NH or alkyl-substituted NH; B = (un)substituted
    ring; D = (un)substituted Ph, pyridyl, pyrazolyl, etc.; R1 = amino,
    formyl, cyanoalkenyl, etc.; R2, R3 = H, (un)substituted alkyl, alkenyl,
    cycloalkyl, etc.; n = 0, 1-4; p, q = 0 or 1] and I salts, stereoisomers
    and tautomers are prepared as acaricides and insecticides.
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    RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological
    study); PREP (Preparation); USES (Uses)
        (preparation as acaricide and insecticide)
    ANSWER 14 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:902883 CAPLUS
DOCUMENT NUMBER:
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143:229846

TITLE: Preparation of anthranilamides as pesticides

INVENTOR(S): Koyanagi, Toru; Morita, Masayuki; Nakamoto, Kenichi;

Hisamatsu, Akihiro

PATENT ASSIGNEE(S): Ishihara Sangyo Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 52 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

APPLICATION NO. PATENT NO. KIND DATE DATE \_\_\_\_\_

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             NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
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                                            WO 2005-JP2351
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                        MARPAT 143:229846
OTHER SOURCE(S):
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$$(R^{1})_{m}$$

$$(R^{1})_{m}$$

$$(R^{2})_{m}$$

$$(R^{3})_{q}$$

$$(R^{3})_{q}$$

AB The title anthranilamides, i.e. N-(2-aminocarbonylphenyl)-1-(2-pyridyl)-1-H-pyrazole-5-carboxamide derivs. represented by the general formula (I) or salts thereof [wherein R1 = halogeno, alkyl, haloalkyl, alkenyl, haloalkenyl, alkynyl, haloalkynyl, alkoxy, haloalkoxy, alkylcarbonyl, haloalkylcarbonyl, alkoxycarbonyl, haloalkoxycarbonyl, (un)substituted phenoxycarbonyl, NO2, CHO; R2, R3 = halogeno, alkyl, haloalkyl, alkoxy, haloalkoxy, cyano; A = Y-substituted alkyl (Y = C3-4 cycloalkyl optionally substituted by ≥1 groups selected from halogeno, alkyl, and haloalkyl); n = 0,1; q = 0-4; provided that R1 is F, C1, Br, or Me substituted at 2-position of the benzene ring and another R1 is halogeno substituted at 4-position of the benzene ring, the 4-halogeno group is F or C1] are prepared They are useful as pesticides, in particular insecticides, acaricides, nematocides, and parasiticides. Thus, 1.49 g

Et3N was slowly added dropwise to a solution of 0.8 g cyclopropylmethylamine hydrochloride in 40 mL THF, stirred at room temperature for 30 min, slowly treated dropwise with a solution of 1 g 2-[1-(3-chloro-2-pyridyl)-3-(trifluoromethyl)-1H-pyrazol-5-yl]-8-methyl-4H-3,1-benzoxazin-4-one in 10 mL THF, and refluxed for 4 h to give, after workup and silica gel chromatoq., 0.54 q N-[6-[[(cyclopropylmethyl)amino]carbonyl]-2methylphenyl]-1-(3-chloro-2-pyridyl)-3-(trifluoromethyl)-1H-pyrazole-5carboxamide (II). II at 3.1 ppm controlled 2-nd to 3-rd instar larvae of Spodoptera litura on cabbage leaves.

REFERENCE COUNT: THERE ARE 62 CITED REFERENCES AVAILABLE FOR THIS 62 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

862995-50-4P 862995-51-5P 862995-52-6P ΤT 862995-53-7P 862995-54-8P 862995-55-9P 862995-56-0P 862995-57-1P 862995-58-2P 862995-59-3P 862995-60-6P 862995-61-7P 862995-62-8P 862995-63-9P 862995-64-0P 862995-65-1P 862995-66-2P 862995-67-3P 862995-68-4P 862995-69-5P 862995-70-8P 862995-71-9P 862995-72-0P 862995-73-1P 862995-74-2P 862995-75-3P 862995-76-4P 862995-77-5P 862995-78-6P 862995-79-7P 862995-80-0P 862995-81-1P 862995-82-2P 862995-83-3P 862995-84-4P 862995-85-5P 862995-86-6P 862995-87-7P 862995-88-8P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of anthranilamides as pesticides such as insecticides, acaricides, nematocides, and parasiticides)

ANSWER 15 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:648522 CAPLUS

DOCUMENT NUMBER: 141:190786

TITLE: Preparation of cyano anthranilamide insecticides Hughes, Kenneth Andrew; Lahm, George Philip; Selby, INVENTOR(S):

Thomas Paul; Stevenson, Thomas Martin E.I. Du Pont De Nemours and Company, USA

PATENT ASSIGNEE(S): SOURCE: PCT Int. Appl., 63 pp.

CODEN: PIXXD2 DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	FENT	NO.			KIN	D	DATE			APPL	ICAT	ION :	NO.		D	ATE	
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		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KP,	KR,	KΖ,	LC,
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			WO	2004-US3568	W	20040121
			KR	2005-700059	A3	20050103
			US	2005-540966	A3	20050629

OTHER SOURCE(S): MARPAT 141:190786

$$R^2$$
 $R^3$ 
 $R^4$ 
 $R^5$ 
 $R^7$ 

The title compds. [I; R1 = Me, Cl, Br, F; R2 = F, Cl, Br, haloalkyl or haloalkoxy; R3 = F, Cl, Br; R4 = H, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, each optionally substituted with one substituent selected from the group consisting of halo, CN, SMe S(O)Me, S(O)2Me and OMe; R5 = H, Me; R6 = H, F, Cl; R7 = H, F, Cl], useful for controlling an invertebrate pest, were prepared E.g., a multi-step synthesis of compound I [R1 = Me; R2 = CF3; R3 = Cl; R4, R5 = H], was given. The compds. I were tested in various biol. tests (data given). This invention also pertains to a composition for controlling an invertebrate pest comprising a biol. effective amount of a compound I, an N-oxide thereof or a suitable salt of the compound I and at least one addnl. component selected from the group consisting of a surfactant, a solid diluent and a liquid diluent.

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RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of cyano anthranilamide insecticides)